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On the fundamentals of organic mixed ionic/electronic conductors

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The first Telluride Science meeting (formerly TSRC) on organic mixed ionic and electronic conductors (OMIECs), Oct 3–7, 2022, brought together researchers across the field to understand the fundamental processes and identify out-standing questions related to this exciting class of materials. OMIECs are organic materials that promote the transport of mobile electronic charge carriers while simultaneously supporting ionic transport and ionic–electronic coupling. These properties open up broad areas of applications from energy to bioelectronics. Devices include batteries, supercapacitors, actuators, electrochromic displays, and organic electrochemical transistors (OECTs). They possess the key strengths of traditional organic electronic materials, such as synthetic tunability and low-temperature processing. Despite the recent advances in devices and applications achieved with such materials, many challenges and gaps in understanding remain. These topics hold the key to designing next-generation materials and devices that continue to push the limits of performance and stability and facilitate novel functionality. This perspective aims to summarize the current understanding, conversations, and debates that made this TSRC particularly engaging, enabling new directions and searching for missing pieces of the OMIEC puzzle.

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The materials – search for model systems

As in other fields within materials science, it is important to identify model systems that will enable accessible and repeatable fundamental studies. At their most basic level, OMIECs support electronic transport through an electronic transport pathway (*i.e.*, along and between conjugated polymer backbones or between nearby redox centers), and ionic transport

occurs through free volume or an ion transport pathway. The ions are needed to ensure charge neutrality when the conjugated or redox system is doped (chemically or electrochemically).

The opportunity and the challenge with organic systems is that such an arrangement can be achieved through many routes: blends, composites, single component systems with varied side chains, block co-polymers, *etc.* A single model system has eluded the field as a whole, and different questions demand different model systems. Poly(3-hexylthiophene)

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(P3HT) is the most studied system in this area. Many fundamental studies have focused on electrochemistry in organic electrolytes, such as TBAPF₆ in acetonitrile. Controlled electrochemical doping led to highly conducting films.¹ P3HT, in combination with salts (*e.g.*, LiClO₄) and poly(ethyleneoxide) (PEO), finds an audience in energy applications or other areas where dry mixed conduction is sought.²

Poly(ethylenedioxythiophene):poly(styrenesulfonate) (PEDOT:PSS) is the most studied OMIEC, a combination of a doped conducting polymer and a polyelectrolyte, and can be regarded as a polyelectrolyte complex. PEDOT:PSS with aqueous electrolytes, including NaCl or KCl, has been well studied owing to their commercial availability and ubiquity in the community of researchers interested in aqueous operation or the role of hydration, *e.g.*, for OECTs or actuators. More recently, conjugated backbones with ionic side groups (so-called conjugated polyelectrolytes) and oligoether side chains fill yet another need and allow operation in aqueous electrolytes. Additional materials strategies from side-chain free polymers and radical polymers are also gaining popularity.

Even so, single component systems are still largely represented by a few chemical structures. While polythiophene backbones are the p-type work-horse of the community, naphthalene diimide (NDI) acceptor units combined with thiophene donors can be regarded as the n-type model system. The electrochemistry of this polymer (*e.g.*, NDI-T) is quite complex but points to a backbone with localized charges on the NDI units.³ Another n-type mixed conductor, the ladder-type poly(benzimidazobenzophenanthroline) (BBL), stands as one of the few side-chain free mixed conductors and has shown high performance in OECTs. Pure p-type redox polymers bear,

for example, triarylamine or carbazole redox units.⁴ On the small molecules front, the films of oligoether tethered fullerenes,⁵ thiophene-flanked diketopyrrolopyrrole units,⁶ polycyclic systems consisting of naphthalene bis-isatin dimer core with rhodanine terminals,⁷ or thiophene-EDOT based cores⁸ are among the structures that can be electrochemically doped and de-doped in aqueous media, while they are yet to be commonly used by different research teams. All these systems can be modified with additives, dopants, and through processing and post-processing.

No one system can answer all the key questions, and no one system is completely ideal (*i.e.*, using PEDOT:PSS in commercial formulations means some additives may not be known to the researcher, and small batch novel synthesized polymers can have variable levels of impurities, defects, and distributions of molecular weights). Nevertheless, some parallels and generalities can be gleaned, putting these diverse materials on a broader spectrum of understanding that benefits the community at large.

The operating environment dictates properties

The composition, structure, and properties of OMIECs are complex and interrelated. Compositionally, all species play a critical role. Structurally, OMIECs often form semi- or *para*-crystalline domains in an amorphous matrix. Multicomponent OMIECs can undergo phase separation, such as the segregation into conjugated polymer-rich and polyelectrolyte-rich domains



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in PEDOT:PSS.^{9,10} Ions and additives affect the degree of crystallinity, coherence length within the crystalline domains, phase separation, and phase purity, amongst other structural parameters.¹¹ Harsh processing techniques can enhance crystallinity and increase the ratio of conjugated polymer to polyelectrolyte in multicomponent OMIECs.¹²

OMIECs have applications in a variety of operating environments. The permeable nature of OMIECs makes them sensitive to operating environments that often include contact with, or immersion in, electrolytes when operated in electrochemical devices. Here again, all species play a critical role, and the role of ion and solvent transport between the OMIEC and its environment must be considered. Contacting electrolytes (*e.g.*, solid polymer, aqueous, organic, or ionic liquid) interact with OMIECs in different ways and to different degrees, depending on characteristics of the solvent (polarity, miscibility) and the ion (size, solvation, polarizability).

The passive swelling (just due to contact with the electrolyte, no electrochemical potential applied) and active swelling (driven by electrochemical potential) of OMIECs must be considered to understand the morphology of the film during operation. During passive swelling, ions and solvent penetrate and swell the OMIEC. The choice of water *versus* acetonitrile electrolytes leads to different degrees of swelling, *e.g.*, determined by *in situ* spectroscopic ellipsometry coupled with cyclic voltammetry.^{13,14} The degree of swelling strongly depends on the miscibility of the electrolyte and the OMIEC, which is often determined by the OMIEC side chain chemistry (oligoether, alkyl, tethered ion) and architecture (length, branching).^{15–18} The miscibility of solvent and ions in the OMIEC can vary, resulting in differential solvent and ion uptake.¹⁹ Further, solvated ions may shed or retain solvation shells when injected into OMIEC. The degree of swelling may differ between amorphous and crystalline domains²⁰ or between conjugated polymer-rich and polyelectrolyte-rich phases.²¹ Ion and solvent uptake into the crystalline domains expands and modifies the crystal structure, in some cases enhancing or disrupting crystallinity dependent on both the OMIEC, electrolyte, and electronic charge density.^{20,22–27}

If the as-prepared OMIEC contains mobile ions that differ from those in the contacting electrolyte, ion exchange will occur, such as ion-proton exchange in PEDOT:PSS.²¹ In some systems, larger concentrations of fixed ionic (polyelectrolyte) charge in the OMIEC introduces Donnan exclusion, limiting ion uptake to oppositely charged counterions.²¹ In such cases, di- and poly-valent counterions can chelate with the fixed charge groups, becoming trapped and inducing ionic crosslinks.²⁸ But in most high-performing OMIECs, Donnan exclusion is suppressed or absent, and cation and anion transport, along with solvent transport, contribute to OMIEC swelling.^{21,29}

A change in the electrochemical potential of the OMIEC can actively modulate how much it swells. The electrochemical oxidation or reduction of the OMIEC modulates the electronic carrier and the counterbalancing ion (dopant) density. This accumulated charge, in turn, can modulate optoelectronic properties – which is core to electroactive functionality of

OMIECs in a variety of applications, *e.g.* electrochemical displays but also electrochemically switchable nanoantennas as metasurfaces.³⁰ Dopant modulation can occur by both ion uptake and expulsion.^{14,29} This modulation of electrostatics also changes the miscibility of the OMIEC and electrolyte surroundings, bringing in or expelling solvent and charge-balanced populations of anions and cations, beyond just the dopant transport.²¹

The degree of active swelling can vary greatly depending on the electrolyte choice and the OMIEC chemical structure. Microscopically, electrochemical cycling can induce crystallite expansion and contraction.^{20,24,25} Extreme active swelling can lead to dissolution/delamination,³¹ but can also be leveraged in reversible swelling/gel formation.³² The effects of active swelling on composition, structure, and properties are often not fully reversible.³³

Additionally, under operating conditions, impurities can play a large role. Dissolved oxygen can limit the electrochemical stability window and accelerate OMIEC degradation.^{34,35} Further, OMIECs with shallow HOMO levels can be spontaneously doped (oxidized).³⁶ Residual catalysts within the OMIEC can determine its electronic conductivity.³⁷ The contacting electrodes also affect the OMIEC, with electrode material (along with oxygen and potential) dictating the potential stability window and degradation kinetics of the OMIEC.³⁴

All this demands the use of multimodal *in situ* characterization in various operating conditions to capture and interrogate the complexities. Structure, composition, and properties must be mapped across relevant environments and potentials. This requires simultaneous (or parallel) probing of electronic (stored charge, electrical conductivity), compositional (individual anions, cations, and solvent), structural (micro-, mesoscale, macroscopic), spectral, and mechanical properties of OMIECs in application-relevant electrolyte environments under controlled potentials. Some popular *in situ* characterization tools are electrochemical UV-VIS-IR absorption spectroscopy, quartz crystal microbalance with dissipation monitoring (QCM-D), X-ray characterization, X-ray photoelectron spectroscopy (XPS), spectroscopic ellipsometry, and scan-probe microscopies.³⁸

Where are the ions in the structure?

A large effort has been made to understand where ions reside within the volume of OMIECs during electrochemical doping/dedoping. However, getting a holistic picture of ions has proven challenging due to the heterogeneous microstructure of OMIECs and the experimental challenges of working with (especially aqueous) electrolytes *in situ*. To understand ion arrangement in OMIECs, the driving force for ion intercalation has to be considered, favourable microstructures have to be found that feature efficient ionic–electronic coupling, and finally, it has to be understood how ions are distributed at device length scales where electric fields can drive their motion.

The ability for ions to penetrate OMIECs is highly dependent on the mixing enthalpy.³⁹ It has been shown that polar



conduction. However, recent work showed that BBL, not only exhibits facile mixed conduction capability, but outperforms the widely studied n-type NDI-T2 based polymers, bearing hydrophilic side chains.⁶⁸ Tang *et al.* showed that BBL is not unique in its ability to exhibit mixed conduction in the absence of solvating side chains. The side-chain free poly(benzodifurandione) synthesized *via* oxidative polymerization in the presence of quinone oxidants appears to be a candidate for bridging the performance gap between n-type and p-type mixed conductors.⁶⁹

What remains evident is that predicting mixed conductor performance based on chemical structure alone is challenging. To understand and utilize the underlying process, it is vital to venture out of what is conventionally known (and assumed) and explore every aspect of polymer chemistry that facilitates and governs mixed conduction.

The role of molecular weight and dispersity

Another outstanding challenge in OMIECs is controlling their molecular weight and dispersity (\mathcal{D} and shape of the molecular distribution), and understanding how these critical molecular parameters influence performance in devices. In semiconducting polymers for OFETs, it is well established, particularly for P3HT, that the molecular weight and dispersity strongly influence the charge mobility. High molecular weight polymers generally have higher mobility due to an increased fraction of tie chains.⁷⁰ It was shown that even a small fraction of low molecular weight polymers in polydisperse samples resulted in a drop in mobility.⁷¹ On the other hand, for mixed transport, the relationship between molecular weight and dispersity, and performance is yet to be established and is likely to be polymer-specific. For example, Wu *et al.* found that higher molecular weight BBL led to increased performance in OECTs.⁷² As described above, the OMIEC system includes the salt – thus, the identity of the electrolyte also affects trends related to molecular weight.²⁰ In multi-component systems, such as PEDOT:PSS, the molecular weight of PSS did not affect the OECT performance, but having a higher \mathcal{D} led to a 4-fold increase in $[\mu C^*]$.⁷³ For establishing clear structure–property relationships for new OMIECs, it will, therefore, be crucial to compare polymers with similar molecular weights and dispersity. This task, however, is likely to be complicated by synthetic challenges, including differences in solubility between the polymers, difficulty in reproducibly synthesizing π -conjugated polymers with controlled molecular weight and dispersity, a similar defect distribution, and variable degrees of purity (*e.g.*, residual catalyst content³⁷).

Post-processing and additives

A different way to modulate mixed conduction that does not require synthetic means is through post-processing. These approaches include molecular doping, solvent blending, and temperature annealing. These are well established for conventional organic semiconductors and their electronic devices, such as OFETs, organic photovoltaics and thermoelectric

generators but have largely been unexplored for OMIECs and OECTs, in particular.

Molecular doping strategies have been applied to OMIECs, both for p- and n-type materials, which have been successful in modulating current densities, shifting threshold voltages, minimizing side reactions, and improving long-term stability in OECTs.^{74–76} Other small molecule additives have been employed to improve the output characteristics of OECTs.⁷⁷ Moreover, the de-doping of PEDOT:PSS has been achieved by promoting interactions between amine groups and PSS-, allowing the generate a polymer for use in enhancement mode OECTs.⁷⁸ While progress has been done in this regard, little is understood about dopant-water interactions (or more generally dopant-solvent interactions), how the dopant interacts with the material itself, and how this extra molecule interacts with the ions in the electrolyte.

On the other hand, solvent blending has been more widely utilized, with co-solvents added to PEDOT:PSS dispersions to improve the conductivity through a change in morphology and.¹¹ The co-solvent approach has also been successful in improving performance of OECTs bearing hole- or electron-conducting OMIECs as the active material. For example, acetone was used as a “bad solvent” to induce aggregation in an n-type NDI-T2 copolymer.⁷⁹ Chlorobenzene was added to a p-type T2-T copolymer to improve the morphology by extending the drying time of the film after spin coating.⁷⁵ Both examples use chloroform as base solvent, which solubilizes the backbone of the polymers better than the additive solvent. Choosing to solubilize either the side chains or the backbone has very different effects on device performance. Solubilizing side chains with polar solvents increases backbone aggregation, which improves the OECT mobility.⁸⁰ This effect is observed not only in polymers but also in small molecule semiconductors.⁸¹

Another approach to improve device performance is *via* an increase of charge density, which can be achieved by chemical dopants. NDI-T2 based polymers were admixed with a Lewis acid, ammonium salt tetra-*n*-butylammonium fluoride, and the additive acted as a simultaneous molecular dopant and morphology-additive, with the combined effects significantly enhancing OECT performance.⁷⁴ Organic salts added to the polymer casting solution, with no energetically plausible mechanism of charge transfer to the OMIEC, improved the backbone planarity and charge delocalization by coupling to the backbone.⁸² The salt-added films show a distinctly porous morphology where the interconnectivity is affected by the salt type and responsible for faster OECT operation.

Another additive type that modulates mixed conduction is ion transporting materials such as cellulose nanofibrils. The ion conductors improved the ion mobility of, *e.g.*, PEDOT:PSS⁸³ and a polythiophene with oligoether side chains.⁸⁴ Besides conductivity enhancing ability, the nanocellulose can act as a reinforcing agent, leading to mechanically adaptive materials that show a reversible decrease in stiffness upon swelling.

The effect of temperature is known to improve crystallinity in organic semiconductors, which in turn can improve field effect mobilities. For OMIECs, for which electrolyte-carrying



ions are needed for conductivity modulation, thermal annealing has so far been considered detrimental, as shown by Flagg, *et al.* in a p-type polymer. By improving crystallinity, the resistance of the active layer to ion penetration is increased. This leads to a lower OECT performance by decreasing mobility.⁶³ However, a recent report showed that operation with a large anion (PF₆ or TFSI) is possible after annealing the film up to 300 °C.⁸⁵ The positive or negative effects of annealing differ from one backbone to another, and seem to be directly tied to their glass transition and melting temperature, rather than to the nature of the side chains themselves.

Many devices comprise OMIEC thin films (on a substrate) with a thickness of typically less than 100 nanometers, which ensures that ions can rapidly reach the complete volume of the material, essential for, *e.g.*, achieving a high switching speed. For some devices such as fiber based OECTs, the material (*e.g.* highly aligned PEDOT:PSS microfibers⁸⁶) may act as both the OMIEC as well as the *de facto* substrate and must hence feature adequate mechanical robustness. The OMIEC material then tends to have dimensions of at least several micrometers, which considerably reduces the speed of ion ingress, but could be mitigated through the use of porous materials.⁸⁷

Emergent phenomena owing to high charge densities

OMIECs display a finite potential window of high electrical conductivity, with the conductance decreasing dramatically beyond a critical gate voltage-controlled charge carrier density. While this phenomenon is reversible and not due to chemical or electrochemical degradation of the OMIEC's π -system, the precise cause is not fully understood yet.

Measurements of *in situ* conductance coupled with cyclic voltammetry show bell-shaped conductance profiles for redox polymer systems.⁴ For P3HT in TBAPF₆/acetonitrile, for example, a plateau-like conductance is found over a large electrochemical potential range of 0.4 up to 0.9 V vs. Fc/Fc⁺ where polaron and bipolaron species are formed.¹ A possible explanation for the behavior may be associated with band filling,^{88–90} while others have proposed that energetic/structural disorder at high carrier densities leads to decreased charge carrier mobility.^{1,91–93} For a further discussion on plateau-like conductivity and the decrease at very high potentials, we refer to electrochemical literature.⁹⁴

In BBL, the maximum electrical conductivity coincides with a change in the polarity of the Seebeck coefficient as the carrier density approaches 1 electron/repeating unit.⁹⁵ This is explained by the formation of multiply charged species and the opening of a hard Coulomb gap around the Fermi energy caused by double occupation and Coulomb interactions occurring at high doping levels. It was also found that the soft Coulomb gap previously proposed to explain the drop in electrical conductivity is of minor importance,⁹⁶ particularly for a rigid polymer like BBL.

Recent work has also shown that the decrease in conductance at high charge carrier density is associated with electron–

cation interactions occurring at the electrolyte and organic semiconductor interface that alter the potential energy landscape in which electronic charges move.⁹⁷ In fullerene (C₆₀) single crystals gated through ionic liquids, the degree of conductance suppression is considerably affected by the size of the ions. Small ions support activated transport and a complete conductance suppression at around 1 electron per C₆₀, while larger ions decrease the extent of suppression and allow exploring transport up to 1.7 s per C₆₀.

Although the decrease in electrical conductivity may pose challenges for conventional logic gates and sensors, it also leads to a unique transistor current response with a Gaussian shape. This response can simulate the activation and deactivation of sodium channels present in biological neurons, creating conductance-based organic electrochemical neurons that accurately replicate crucial biological neural features.⁹⁸ Therefore, these neurons can serve as event-based neuromorphic sensing and processing components, displaying exceptional biorealistic capabilities.

Modeling and simulation

Both quantum chemistry and molecular dynamics (MD) simulations have been used to interrogate the transport mechanisms and structure–function relationships of OMIECs. Recent work⁶¹ shows how grazing-incidence resonant X-ray diffraction (GIRXRD) complemented by periodic DFT calculations allows the determination of the mean position of the ion relative to the polymer backbone in the crystalline regions, which somewhat surprisingly is found near the lamellae midpoint (and not close to the polymer backbone). Multiscale methods combining MD and Kinetic Monte Carlo (KMC) have been instrumental as a tool to describe charge transport and percolation in conjugated materials and will continue to be key in OMIEC research. A multiscale transport model based on information from the structural and electrical characterization of PEDOT:PSS blends was also recently reported to explain the exponential-like conductivity scaling at different PEDOT:PSS ratios.⁹⁹ Charge transport from one PEDOT-rich grain to the next happens thanks to isolated/dispersed PEDOT chains in a PSS-rich matrix, with PSS-conjugated groups enhancing tunneling efficiency.

A coarse-grained multiscale model was used to systematically investigate the effects of side chain polarity in a series of random OMIEC copolymers.¹⁹ The highest charge mobility (and doping) is associated with the highest fraction of polar side chains. The results where a similar fraction of polar and apolar side-chains were mixed point to a significant drop in charge mobility due to a shorter polaron delocalization length arising from the increased torsional disorder along the polymer backbone.

Multiscale models also shed light on time-dependent ion-charge interactions in mixed-conducting materials. Recent work¹⁰⁰ has incorporated DFT calculations and MD simulations to capture the rapid charge fluctuation on the polymer backbone due to the presence of solvent and ions. While this



approach does not consider charge transfer across polymer chains, it captures both ion-induced instantaneous polarization and slow (~ 1 ns) charge reorganization due to conformational changes in polymer chains. Interactions across polymer chains dominate charge dynamics, which are expected to play a major role in bulk polymer aggregates or crystallites. While promising, this study highlights how representing the interaction between ionic and electronic charge carriers remains the most challenging task for any computational model of OMIECs. Although some methodological barriers still exist for reliably reproducing amorphous and semi-crystalline morphologies and ion-polaron coupling, ample opportunities exist for using already mature methodologies to study other structure-function and morphology-function relationships in OMIECs. Such efforts are made urgent by the extreme complexity and potential variety of OMIEC materials (ionic species, polymer backbone and connectivity, side chain polarity, length, and frequency). Despite the number of unknowns in the field, we believe a systematic OMIEC optimization effort guided by computational screening will soon be within reach as the community gradually uncovers relationships and correlations.

Spanning different systems

Despite some generalities across systems and common tools to study them, different systems and applications are governed by very different phenomena. For example, electronic transport is mechanistically different in non-conjugated *versus* conjugated redox polymers. Specifically for non-conjugated redox polymers, the electronic charge is localized on pendant groups where electrons can hop from site to site.¹⁰¹ In contrast, conjugated polymers distribute charge through the polymer's pi-conjugated backbone.

For electron conduction in non-conjugated redox polymers and small radical molecules, it is important to consider the proximity of the redox sites. As the site-to-site distance decreases, the conductivity increases. For example, a 1000-fold difference in the solid-state conductivity occurs when changing the hopping distance from 5.7 to 11.6 Å in certain crystals of small radical molecules.¹⁰² For non-conjugated redox polymers, the flexibility of the polymer chain must be considered in addition to redox site proximity.¹⁰³ As the chain's flexibility increases, the diffusivity of the polymer increases, and there are more opportunities for redox sites to come within closer proximity through thermal motion. For example, the highly flexible and low- T_g radical-containing polymer poly(4-glycididloxy-2,2,6,6-tetramethylpiperidine-1-oxyl) exhibited the highest solid-state conductivity within its class of 28 S m⁻¹. In the case of carbazole redox polymers (polyvinylcarbazole), conductivities as high as 2×10^{-2} S cm⁻¹ could be obtained upon chemical doping.⁴ In this case, the simultaneous cross-linking and doping of the starting materials into bicarbazole redox dimers could be proven by a combination of UV-vis-NIR spectroscopy and *in situ* and *ex situ* conductivity measurements.

In contrast to dry conditions, non-conjugated redox polymers exhibit mixed ion-electron-solvent transport in electrochemical

cells. The charge transport is affected by polymer-solvent and polymer-ion interactions, swelling, and the ion's solvation shell.¹⁰⁴⁻¹⁰⁶ For example, EQCM-D experiments revealed that both anions and cations could participate in the polymer's redox behavior, which should be considered in future designs.¹⁰⁴ Further, electrolyte choice can yield 1000 \times changes in the charge storage of non-conjugated redox polymers,¹⁰⁵ indicating a rich future design space for electrolytes.

The combination of ac-impedance spectroscopy and dc measurements of mixed conductors (PEDOT:PSS and conjugated polyelectrolytes) can help to decouple electronic and ionic charge transport as a function of relative humidity and corresponding water uptake.^{15,107} The influence of relative humidity was particularly prominent for the ion conductivity, which reached 10⁻² S cm⁻¹ in the wet state. The electronic conductivity is less sensitive to humidity but requires the existence of electronic percolation pathways throughout the films, which is directly interrelated with the morphology upon exposure to humidity. It is interesting to note that the humidity. Response of the polyelectrolyte phases could be used to tune mechanical properties and make actuators.¹⁰⁸

When translating OMIECs into dry systems, the advantages conferred by solvent-assisted ion transport are compromised, and instead, the segmental mobility of the ion-conducting chain becomes a determining factor in the intrinsic limitations of ion transport. In the case of side-chain-based OMIECs, it becomes essential to carefully consider the structure of the side chains, ensuring optimal ion solvation and promoting fast dynamics to facilitate ion motion. Recent joint computational and experimental studies have provided valuable insights into the intricacies of Li-salt-doped OMIECs based on oligoethylene glycol-substituted thiophene derivatives.¹⁰⁹⁻¹¹² These reports on dry OMIECs highlight the challenge of effectively separating the dynamics of the side chains responsible for ion transport from the more rigid conjugated core imparting semicrystallinity for efficient electronic transport.

The rate at which an OMIEC transports ions is affected by whether or not the ions are contained within a solvent shell, as is typically the case for PEDOT:PSS and oligoether side chain containing polymers in aqueous electrolytes.^{56,63,113,114} In other cases, for example, for P3HT with polymer electrolytes, the ion moves without a solvent shell, where solubility interactions dominate.¹¹⁵ It is typically, but not always, observed that a solvent shell improves ion transport. Nevertheless, a single perspective unifying the two extremes and providing a predictive structure-property relationship remains challenging.

Conclusions and perspectives

While our focus has been on conjugated polymers and organic small molecules, it is clear that the library of OMIECs continues to expand by the day. Small molecules, for example, have found an important place in precision molecular design and the extreme case of poly-crystalline morphologies. MOFs, COFs, and porous conjugated polymers, while their integration into



devices presents barriers currently, afford three-dimensional frameworks where porosity can be controlled at the molecular scale, and chemical functionality can be finely dialed in. As for electrolytes, ionic liquids, as first investigated in electrolyte-gated transistors, are now numerous, and biocompatible versions with various ionic units may introduce new characteristics to the same OMIEC devices. Broadening the library of mixed conductors beyond polymers – perhaps even introducing hybrid systems – will thus open new applications for OMIEC devices.

The quest for model systems is useful but may be futile given the breadth of conditions and target applications under the OMIEC tent. Nevertheless, they prove useful for fundamental studies. The OMIEC must be considered a complete system, including the extrinsic or intrinsic ions, additives, and solvents, as appropriate. In addition, given the volatility of the solvents or the extrinsic nature of some electrolytes, multi-modal and operando characterization is needed to make direct device-material connections. Many differences in molecular design and film microstructures are not considered or well controlled, adding further challenges. Moving forward, questions remain about the exact location, interactions, and compositions (ionic) within OMIEC materials and how to model them properly. Can such understanding unlock designs for ideal materials? More stable formulations? Or allow for less common and highly non-linear responses toward novel applications? It's likely, but the community needs to tackle these questions collaboratively to get there.

Conflicts of interest

There are no conflicts to declare.

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